Paralytic Shellfish Poisoning: The Relationship Between *Alexandrium* Abundance and PSP Toxins on Kodiak Island, Alaska

Julie A. Matweyou and Dean A. Stockwell

Institute of Marine Science

Christopher A. Scholin

Monterey Bay Aquarium Research Institute

Sherwood Hall

Marine Toxin Laboratory, U.S. Food and Drug Administration

Vera L. Trainer

NOAA/NMFS/ECD

Jason D. Ray

Saigene Corporation

Terry E. Whitledge, Amy R. Childers and F. Gerald Plumley

Institute of Marine Science

Abstract

Paralytic Shellfish Poisoning (PSP) has severe negative impacts in Alaska. This study was designed to improve existing PSP monitoring programs by incorporating phytoplankton monitoring. *Alexandrium* abundance was ascertained at a near shore site on Kodiak Island during 2000 and 2001 using species-specific LSU rRNA targeted oligonucleotide probes in whole cell (WC) and sandwich hybridization (SH) assay formats. *Alexandrium* abundance exhibited two distinct peaks (>400 cells/L) in 2000, both of which lasted approximately 2 weeks. *Alexandrium* abundance in 2001 was much more sporadic, with 4-5 peaks, each lasting only 3-7 days. In 2001, *Alexandrium* abundance tracked water column toxicity as determined via a ³H-Saxitoxin receptor-binding assay. Importantly, DNA probe data revealed a correlation between *Alexandrium* abundance and blue mussel (*Mytilus edulis*) toxicity in both 2000 and 2001. The results also demonstrated that increases in *Alexandrium* abundance preceded elevated toxin levels in shellfish suggesting that this method could prove useful as a monitoring tool to predict toxic events prior to shellfish harvest. Overall, this report provides compelling evidence that DNA probe chemistry can be used to estimate the abundance of *Alexandrium* in the field, however a number of problems must be rectified if the assay is to be used for monitoring purposes.